

IO25020

Bayer CropScience



March 26, 2013

Document Processing Desk 6(a)(2)
Office of Pesticide Programs (7504P)
U. S. Environmental Protection Agency
Room S-4900, One Potomac Yard
2777 South Crystal Drive
Arlington, VA 22202-4501

RE: 6(a)(2) Incidents Accumulated for the Month of February 2013

Dear Sir/Madam:

Reportable incidents accumulated for the month of February 2013 for Bayer CropScience and Bayer Environmental Science are attached.

Bayer CropScience
RTP
P. O. Box 12014
RTP, NC 27709
Tel. 919 549-2000

The information with this letter is being submitted to the EPA pursuant to the Agency's interpretation of requirements imposed on registrants by Section 6(a)(2) of FIFRA. This information does not necessarily constitute additional factual information regarding unreasonable adverse effects within the meaning of 6(a)(2). It is being submitted to enable the Agency to make its own assessment of the information.

If you have questions or concerns, please do not hesitate to contact me at any time.

Sincerely,

A handwritten signature in black ink, reading "Gerret Van Duyn".

Gerret Van Duyn
Compliance Manager
State Regulatory and Documentation Services
919-549-2914

CC: AE Coordinator, CA Department of Pesticide Regulation
Jeanine Broughel, NY Department of Environmental Conservation

/attachment



Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area Page 1 of 3

Row 1 Administrative Data	Reporter Name [REDACTED]	Submission date. 3/26/2013	Contact person (if different than reporter)	Internal ID 1126133
	Address [REDACTED]		Address	
	Phone # [REDACTED]		Phone #	
	Incident Status: <i>New</i>	Location and date of incident <i>Charlotte, NC USA 02/25/2013</i>	Date registrant became aware of incident. <i>02/27/2013</i>	Was incident part of larger study? <i>No</i>
Row 2 Pesticide(s) Involved	EPA Registration # (Product 1) <i>72155-80</i>	EPA Registration # (Product 2)		EPA Registration # (Product 3)
	A.I. (s) <i>Beta-Cyfluthrin, sodium o-phenylphenate</i>	A.I. (s)		A.I. (s)
	Product 1 name <i>Home Pest plus Germ Killer Indoor & Outdoor Killer RTU (1 Gal)</i>	Product 2 Name		Product 3 Name
	Exposed to concentrate prior to dilution? <i>No</i>	Exposed to concentrate prior to dilution?		Exposed to concentrate prior to dilution?
	Formulation	Formulation		Formulation
Row 3 Incident Circumstances	Evidence label directions were not followed? <i>No</i> Intentional misuse? <i>No</i>	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/ woods, agricultural (specify crop) right-of- way (rail, utility, highway)). <i>Own Residence</i>		Situation (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/ formulating). <i>See Incident Description Notes</i>
	Applicator certified? <i>UNK</i>			
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) <i>See Incident Description Notes</i>			

Personal privacy information

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Brief description of incident circumstances.

Wilson, Lauren Feb 27 2013 4:01PM

Hx Caller states that she sprayed baseboards and cracks in home 2 days ago. Her kids where playing on the sofa while caller was spraying, so she does not know of any direct exposure. Caller noticed a red mark on child's face, but didn't think much of it at first. That night caller noticed that it started to spread on her face and down to her shoulder/arm. Caller could then tell that it was hives. She called nurse and was instructed to give Benadryl. The Benadryl worked but then the hives came back by morning and it was worse (legs, face, back & stomach). She took to pediatrician. Dx with allergies and recommend continue with Benadryl and Zyrtec. Caller has appointment with Allergist March 7th.

A Recommend continue care with MD. Will document incident. Recommend having MD call for complete ingredient list. Case # provided.

Yeager, Greg Mar 7 2013 1:54PM

Attempted CB. Left a message requesting follow up. Reset.

Marshall, Josephine Mar 7 2013 2:05PM

CB from [REDACTED]. She took her daughter to the allergist today but has not received any results yet. Initially the MD believes that the hives were due to the virus that the child had at that time. The child is now asx and they are waiting for results from blood tests that were done today. No further treatment was done other than the exam and testing. Symptoms cleared up on approximately 03/05/2013.

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Demographic information: Age: 5 Year(s) Sex: Female Occupation (if relevant) NA	Exposure route: Unknown route	Was adverse effect result of suicide/homicide or attempted suicide/homicide? No	Was protective clothing worn (specify)? None Reported
If female, pregnant? NO	Was exposure occupational? Not indicated If yes, days lost due to illness: NA	Time between exposure and onset of symptoms: 30 min or less	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). Private MD/DVM-treated & released	List signs/symptoms/adverse effects Dermatological-Hives/Welts	If lab tests were performed, list test names and results (If available, submit reports) None Reported	
Exposure data: NA Amount of pesticide: NA Exposure duration: Acute < 8hrs Patient weight: Unknown			
Human severity category: HC			
This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)			
			Internal ID # 1126133